Masses of ⁷⁶Kr and ⁷⁴Kr

D. M. Moltz and K. S. Toth Oak Ridge National Laboratory, Oak Ridge, Tennessee 37830

R. E. Tribble, R. E. Neese, and J. P. Sullivan Cyclotron Institute, Texas A&M University, College Station, Texas 77843 (Received 25 June 1982)

The masses of ⁷⁶Kr and ⁷⁴Kr were measured utilizing the ⁷⁸Kr(⁴He, ⁶He) and ⁷⁸Kr(⁴He, ⁸He) reactions. Mass excesses of -68971 ± 13 and -62210 ± 75 keV were obtained for ⁷⁶Kr and ⁷⁴Kr, respectively, and found to compare favorably with some mass formulae predictions, but not with others. Speculations about shape differences in light krypton and rubidium isotopes are presented as they relate to various mass formulae.

NUCLEAR REACTIONS ${}^{4}\text{He} + {}^{78}\text{Kr}$, $E_{\text{lab}} = 79.7$ and 102.7 MeV; measured $E_{6_{\text{He}}}$, $E_{8_{\text{He}}}$; ${}^{76}\text{Kr}$, ${}^{74}\text{Kr}$ deduced Q, masses; magnetic spectrograph. Enriched gas targets.

I. INTRODUCTION

A study of the mass systematics of nuclei far from stability is a sensitive way of comparing nuclear models. It also presents an interesting method for demonstrating specific nuclear phenomena. For instance, model independent two-neutron separation energies (S_{2n}) readily exhibit the standard nuclear shell structure. Sometimes the mass systematics of an A' = A + 4k, $T_z = constant$ series of nuclides yields information about nuclei in a particular mass region. One such inference can be made from the $A = 4N + 1, T_z = +\frac{1}{2}$ series of beta-delayed proton emitters $^{1-3}$ as they relate to shape contributions to the nuclear wave functions between the $f_{7/2}$ and $g_{9/2}$ shells. These nuclei are particularly interesting because of the proximity of the proton-drip and Z = N lines.

A recent foray into this region consisted of direct mass measurements^{4,5} for most of the known rubidium isotopes. Since that time, we have begun a systematic study⁶ of the light neutron-deficient isotopes of rubidium and krypton to determine accurate beta endpoints with an intrinsic germanium detector. (Earlier plastic scintillator measurements have been found to be in error, requiring masses obtained in this manner to be reexamined; the analysis technique used in Ref. 6 has been described⁷⁻⁹ elsewhere.) Nearer the valley of stability, however, many unstable proton-rich nuclei decay only by electron capture. In some of these instances transfer reactions may be utilized to measure nuclear masses. One such pivotal nucleus with a low Q_{β} value whose mass had not been measured is ⁷⁶Kr. We therefore undertook to determine its mass excess by using the ⁷⁸Kr(⁴He,⁶He) reaction. As part of the study we also utilized the ⁷⁸Kr(⁴He,⁸He) reaction to measure the mass of ⁷⁴Kr; its decay energy to ⁷⁴Br is known¹⁰ to ± 120 keV. Since the start of our investigation, the mass of ⁷⁶Kr was measured¹¹ via the ⁷⁸Kr(*p*,*t*) reaction.

II. EXPERIMENTAL PROCEDURE

The mass of ⁷⁶Kr was determined by measuring the Q value for the ⁷⁸Kr(⁴He,⁶He)⁷⁶Kr reaction at $E_{^{4}\text{He}} = 79.7$ MeV; the mass of ⁷⁴Kr was then measured via the ⁷⁸Kr(⁴He,⁸He)⁷⁴Kr reaction at $E_{^{4}\text{He}} = 102.7$ MeV, an incident energy chosen to permit simultaneous observation of ⁶He and ⁸He particles leading to the ground states of ⁷⁶Kr and ⁷⁴Kr. Alpha-particle beams from the Texas A&M University 224-cm cyclotron were incident upon enriched krypton targets (90+% ⁸⁰Kr and 99 +% ⁷⁸Kr) maintained in a gas cell statically pressurized to 40–60 Torr at 300 K. This ~100 cm³ gas cell was isolated from the surrounding vacuum by ~1.9 mg/cm² Havar foil. The well-collimated reaction products were detected on the focal plane

1914



FIG. 1. Gated ⁶He position spectra at $E_{4_{\text{He}}} = 79.7$ MeV for the reactions: (a) ${}^{80}\text{Kr}({}^{4}\text{He},{}^{6}\text{He}){}^{78}\text{Kr}$ and (b) ${}^{78}\text{Kr}({}^{4}\text{He},{}^{6}\text{He}){}^{76}\text{Kr}$. Large arrows indicate detector limits. Peaks labeled by numerals correspond to: (1) ${}^{78}\text{Kr}$ ground state, (2) first 2⁺ excited state in ${}^{78}\text{Kr}$, and (3) ${}^{76}\text{Kr}$ ground state.

of an Enge split-pole magnetic spectrograph by a 10-cm single-wire gas proportional counter, used as a ΔE detector, with a 5.0×1.0 cm×600 μ m solid state silicon detector serving as an *E* detector. The time-of-flight (TOF) of the particles through the spectrograph [relative to the cyclotron radio frequency (rf)] and the particle position obtained via charge division in the gas counter were stored in addition to the *E* and ΔE signals in event-by-event mode on magnetic tape for subsequent playback. Figure 1 shows a typical position spectrum gated by suitable bounds for ⁶He particles in the *E*, ΔE , and

TOF spectra. This experimental setup has been shown¹² to reject spurious background events at a level below 200 pb/(sr MeV) for medium mass targets. Additional experimental details may be found elsewhere.¹³⁻¹⁵

While the experiment consisted of two distinct mass measurements, several parameters remained unchanged throughout. Elastically-scattered 79.7-MeV ⁴He²⁺ particles were used to determine the vertical efficiency of 0.82. The spectrograph was placed at $\theta_{lab} = 8^{\circ}$ with an entrance aperture corresponding to 2.6 msr. The laboratory angle was chosen after comparison with the ⁸He angular distribution¹⁶ arising from the ⁶⁴Ni(⁴He,⁸He)⁶⁰Ni reaction. A 0.2-mm Kapton absorber foil was placed between the gas proportional counter and the silicon detector to ensure that the ⁸He particles would stop in the silicon counter. After initially charging the gas cell with 40 Torr of ⁸⁰Kr, calibration of the position spectrum was accomplished by varying the spectrograph magnetic field setting to change the position of the ⁶He particles associated with the ground state of ⁷⁸Kr. Figure 1(a) represents the final ⁶He spectrum arising from ⁸⁰Kr. The ⁸⁰Kr gas was then removed and was replaced with 40 Torr of ⁷⁸Kr gas resulting in the spectrum shown in Fig. 1(b). All spectra were collected in short runs corresponding to an average beam intensity of $\sim 1.0 \,\mu A$ and an integrated beam of 2.5 mC.

The ⁴He beam energy was then raised to 102.7 MeV and the ⁷⁸Kr target pressure to 60 Torr. The rest of the experiment was conducted under these conditions. The integrated beam for this second part was 74.7 mC at an average beam intensity of $\sim 1.2 \ \mu$ A. On-line data sorting was employed to record the ⁶He rate as a monitor of the relative stability of all experimental systems.

III. RESULTS AND COMPARISONS

The ⁶He position spectrum arising from the ⁸⁰Kr(⁴He,⁶He)⁷⁸Kr reaction is shown at the bottom of Fig. 1, while the top portion of this figure corresponds to a ⁶He spectrum due to the ⁷⁸Kr(⁴He,⁶He)⁷⁶Kr reaction obtained with the same cyclotron and spectrograph parameters. The measured Q value of -20.351 MeV yields a mass excess of -68971 ± 13 keV for ⁷⁶Kr. The 13-keV uncertainty arises from a centroid and calibration uncertainty of 10 keV added in quadrature to the 8-keV uncertainty¹⁷ in the ⁷⁸Kr mass excess. The 324 integrated peak events in Fig. 1 correspond to a laboratory cross section of 6 ± 2 µb/sr for the ground

state of ⁷⁶Kr averaged over the 2.6 msr solid angle centered at $\theta_{lab} = 8^{\circ}$. The width in the observed peak is due primarily to energy straggling in the gas target and the Havar isolation foil.

Raising the bombardment energy to 102.7 MeV insured the simultaneous observation of ⁶He and ⁸He particles at the detector location on the focal plane. Although the ⁶He position spectrum could provide a convenient system check, the primary source of background in the ⁸He spectrum came from ⁷Li events. These ⁷Li particles exhibit almost identical E and TOF spectra to those of ⁸He, but $\Delta E_{\gamma_{\rm Li}} \simeq 2\Delta E_{^{8}_{\rm He}}$, permitting an effective separation and identification. A two-dimensional plot of E vs TOF is shown in Fig. 2. The ΔE gate used here was set by observing a null effect upon the ${}^{4}\text{He}^{+}$ peak. $({}^{4}\text{He}^{+} \text{ and } {}^{8}\text{He}^{2+} \text{ have the same } dE/dx.)$ The ${}^{8}\text{He}$ peak remained intact while the ⁷Li peak decreased by a factor of 50 from the ungated spectrum. The ⁸He gated position spectrum is shown in Fig. 3. The four integrated counts correspond to a laboratory cross section of 1.4+0.8 nb/sr at $\theta_{lab} = 8^\circ$.

The Q value for the 78 Kr(4 He, 8 He) reaction was found to be -41.12 MeV which yields a mass excess of -62210 ± 75 keV for 74 Kr. The 75-keV uncertainty was calculated from a 70-keV peakcentroid and calibration uncertainty and a 30-keV beam energy uncertainty. This latter value is estimated from past measurements with the analyzing magnet contained within the cyclotron beam optics system. Since this value contributes very little to the final error, a direct beam energy measurement was deemed unnecessary. The one higher energy event is consistent with the background observed with a 99 + 20 Ne target used with this gas cell in a new measurement¹⁸ of the mass of the unbound nuclide 16 Ne via the 20 Ne(4 He, 8 He) 16 Ne reaction.

Our value of -68971+13 keV for the mass excess of ⁷⁶Kr agrees well with a recent¹¹ value obtained from the 78 Kr(p,t) 76 Kr reaction of -68968 + 15 keV. This excellent agreement permits us to adopt a value of -68970 ± 11 keV for the ⁷⁶Kr mass excess. The measured cross section for this latter reaction is approximately two orders of magnitude larger than that for the (⁴He,⁶He) reaction. This large discrepancy is somewhat surprising because in principle these reactions may proceed via three differing mechanisms¹⁹: direct two-neutron transfer, indirect multistep two-neutron transfer, and two sequential one-neutron transfers. The sequential transfer might seemingly be more important for the (⁴He, ⁶He) reaction than for the (p,t) case because of the greater similarity between the ⁵He



FIG. 2. Two-dimensional plot of energy (abscissa) vs TOF (ordinate) with a ${}^{4}\text{He}{}^{+}\Delta E$ gate. Labeled peaks correspond to: (a) ${}^{4}\text{He}{}^{+}$, (b) ${}^{8}\text{He}{}^{2+}$, and (c) ${}^{7}\text{Li}{}^{2+}$.

and ⁶He wave functions than for the deuteron and triton wave functions. Jang and Sorensen¹⁹ conclude that this sequential transfer is not negligible for the (⁴He,⁶He) reaction. However, the direct process would tend to favor the enhanced cross section for the (*p*,*t*) reaction over the (⁴He,⁶He) reaction, possibly indicating the dominance of the direct channel in this mass region, an idea which has been previously espoused.²⁰ The cross section for the (⁴He,⁸He) reaction (1.4 ± 0.8 nb/sr) is consistent with other results¹⁶ in this region. Additionally, the ratio ($d\sigma/d\Omega$)_{8He}/($d\sigma/d\Omega$)_{6He} is not extraordinary.



FIG. 3. Gated ⁸He position spectrum at $E_{4_{\text{He}}} = 102.7$ MeV for the ⁷⁸Kr(⁴He,⁸He)⁷⁴Kr reaction. Arrows indicate detector limits.

Since evidence for a similar direct process for the four-neutron transfer in the $({}^{4}\text{He}, {}^{8}\text{He})$ reaction has been presented 16 elsewhere, this could further be interpreted as evidence (albeit indirect) for the notion of a primarily direct process in the $({}^{4}\text{He}, {}^{6}\text{He})$ reaction.

The mass excess of -62210 ± 75 keV for 74 Kr disagrees with the accepted value^{$\overline{17}$} of -62020 ± 100 keV based upon a Q_β mass difference measurement¹⁰ utilizing a plastic scintillator beta spectrometer. Since the mass of $^{74}Br (-65295 \pm 15 \text{ keV})$ was determined accurately by a (p,n) threshold reaction,²¹ this difference probably originates with an incorrect level structure for ⁷⁴Br or the inherently bad energy resolution of the plastic scintillator system. These two problems are not uncommon in beta endpoint measurements (see, e.g., Ref. 6). We therefore adopt our value of -62210 ± 75 keV for the mass excess of 74 Kr. The relative importance of these adjustments to the mass excess values and their uncertainties will be discussed in terms of the overall mass surface in the next section.

IV. DISCUSSIONS AND CONCLUSIONS

It is often necessary to view a particular set of mass measurements as part of a larger systematic picture. This approach, adopted for the remainder of the discussion, is particularly useful in delineating regions of deformation such as that found in the



FIG. 4. Mass excesses plotted vs neutron number for nuclei with Z = 33 - 38 and N = 37 - 42.

neutron deficient krypton, rubidium, and strontium nuclei. Table I lists the updated mass excess values for nuclei with (Z = 33 - 38, N = 37 - 42) while Fig. 4 graphically depicts this same region. Many mass

TABLE I. Partial tabulation of updated mass excesses in keV. Uncertainties are enclosed in parentheses. Numbers given in the lower line represent calculations made in this investigation using the transverse Garvey-Kelson relationship (see text). Values have been taken from Ref. 17 unless otherwise noted.

Z	N = 37	38	39	40	41	42
38(Sr)			-57830(210) ^{a,b}		-65600(130)°	
	-47950(460)	-55165(250)	- 58869(135)	-63360(115)	-65400(85)	-70395(75)
37(Rb)	-51984(270) ^d	-57491(188) ^d	$-60906(45)^{e}$	$-65209(50)^{e}$	$-66908(22)^{\rm f}$	$-70902(42)^{d}$
	-52156(240)	58010(100)	- 59789(100)	-64666(70)		
36(Kr)	-56980(140)	$-62210(75)^{a}$		-68970(11) ^{a,g}	$-70481(43)^{e}$	-74150(8)
	56400(90)	-62455(60)	-63917(17)	69489(50)	-70342(20)	
35(Br)		-63670(220)	-65295(15)	-69159(20)	-70303(15)	-73241.5(38)
	-59169(35)	-63693(21)				
34(Se)		-67894(12)	-68209(11)	-72212.7(26)	-72169.0(24)	-75259.2(25)
	-63346(25)					
33(As)	-64339(20)	-67893(4)	-68232(7)	- 70949(4)	-70859.7(25)	-73033.9(23)

^aThis work.

^bReference 1.

^cReference 22.

^dReference 5.

^eReference 6.

^fReference 23.

^gReference 11.

prediction formulae use known mass excesses to obtain a series of parameters for semiempirical calculations. Unfortunately, these parameters might dramatically change with new input masses. The Garvey-Kelson^{24,25} mass relationship may be used in either this latter way or in a recursive building sense. The transverse form of this equation for neutron-deficient nuclei, written as

$$M(A,T_Z) \simeq M(A,T_Z+2) + M(A+1,T_Z+\frac{1}{2}) - M(A+1,T_Z+\frac{3}{2}) + M(A-1,T_Z+\frac{1}{2}) - M(A-1,T_Z+\frac{3}{2}),$$
(1)

can be solved as a partial differential equation for $M(A, T_Z)$. (This approach is utilized by Jänecke.²⁶) Another approach utilizes the recursive nature of the formula to build newer predictions further from stability. This method of course requires good input masses to yield reliable mass predictions. The second (or lower) entry for many of the nuclei listed in Table I is the continued reapplication of this relationship. Quoted uncertainties represent only the accumulated errors from input values.

One of the most instructive and informative ways to view new mass values on the nuclidic mass surface is through comparisons with differing theoreti-

cal models, including the revised Garvey-Kelson method delineated earlier. Table II lists predictions from several models and the differences between them and the masses obtained in this work and other recent results⁶ in this region. Myers's droplet model²⁷ consistently overpredicts the stability of these masses, as is indicated by the similarity of the average $(\langle \Delta \rangle)$ and the root-mean-square average $(\Delta_{\rm rms})$. This case may be contrasted with the revised Garvey-Kelson predictions where $|\langle \Delta \rangle| \neq |\Delta_{\rm rms}|$. This situation is predicated upon the ⁷⁷Sr case which requires the mass of ⁷⁶Rb (a mass which only Groote et al.²⁷ predict with even relative success). Thus, all of the Garvey-Kelson based predictions give similar results. Particular attention should be given to the excellent agreement between the experimental ⁷⁶Kr mass and the prediction of Möller and Nix.²⁸ The primary idea behind their calculations²⁹ incorporates shape independent and dependent effects and requires the empirical determination of only five constants by fitting known masses. Thus, one might expect that the nuclidic mass surface calculated in this manner would be sensitive to any shape changes. The mass of any predicted deformed nuclei would provide some small test for this hypothesis. The predicted onset of prolate deformation in the krypton isotopes is at ⁷⁶Kr. Unfortunately, this deformation onset in the rubidium

TABLE II. Experimental mass excesses (in MeV) for ⁷⁴Kr, ^{76,77}Kr, ^{76,77}Rb, and ⁷⁷Sr compared with predictions from various mass models.

Isotope	Experimental mass	Revised Garvey-Kelson	Jänecke ^a Garvey-Kelson	Möller ^b and Nix	Myers ^a	Liran ^a and Zeldes	Comay ^a and Kelson	Groote, Hilf, ^a and Takahashi			
⁷⁴ Kr	-62.210(75)	-62.46	-62.32	-62.09	- 64.04	-62.20	-62.45	-62.66			
⁷⁶ Kr	-68.970(11)	69.49	-69.07	-68.98	-70.22	-68.91	-69.16	-68.89			
⁷⁷ Kr	-70.481(43) ^c	-70.34	-70.12	-69.58	-71.35	-69.80	- 70.19	-70.26			
⁷⁶ Rb	-60.906(45) ^c	- 59.79	- 59.98	-60.23	-62.05	- 59.21	- 59.94	- 60.66			
⁷⁷ Rb	$-65.209(50)^{\circ}$	-64.77	-64.91	-65.04	-66.54	-64.12	-64.91	-65.25			
⁷⁷ Sr	- 57.830(210) ^d	- 58.87	- 57.30	- 57.77	- 59.67	-56.21	-57.53	-58.14			
			Difference (experiment minus theory)								
⁷⁴ Kr		0.25	0.11	-0.12	1.83	0.01	0.24	0.45			
⁷⁶ Kr		0.52	0.10	0.01	1.25	-0.06	0.19	-0.08			
⁷⁷ Kr		-0.14	-0.36	-0.90	0.87	-0.68	-0.29	-0.22			
⁷⁶ Rb		-1.12	-0.93	-0.68	1.14	-1.70	-0.97	-0.25			
⁷⁷ Rb		-0.44	-0.30	-0.17	1.33	-1.09	-0.30	-0.04			
⁷⁷ Sr		1.04	-0.53	-0.06	1.84	-1.62	-0.30	-0.31			
	$\langle \Delta \rangle$:	0.02	-0.32	-0.32	1.38	-0.86	-0.24	0.04			
	$\Delta_{\rm rms}$:	0.69	0.48	0.47	1.42	1.09	0.47	0.26			

^aReference 27.

^bReference 28.

^cReference 6.

^dReference 1 and this work.



FIG. 5. Mass formulae comparisons for krypton and rubidium isotopes with N = 37 - 43. Points marked with an imes indicate values obtained via the revised Garvey-Kelson method given in the text.

isotopes occurs around $A \sim 80$, a member of the original ensemble of averaged masses. A systematic study of the strontium mass excesses would therefore be of great value for comparison.

Figures 5 and 6 show graphical representations of the differences between the experimental masses and several theoretical predictions for the krypton and rubidium isotopes. Generally, all of the predictions incorrectly give the mass of ⁷⁶Rb. Although the value of Groote et al.²⁷ is not terribly different, a definite trend reversal is evident. If there exists some quasimagic number around N = 39 in these deformed nuclei, then the mass of ⁷⁵Kr must be obtained in addition to improving the error bar for ⁷⁷Sr. Two-neutron separation energies in this region could be interpreted as supporting this claim; however, error bars on several entries are far too large to generalize in this manner. The mass of ⁷⁵Kr should also provide an additional clue in the even-Z neutron pairing energy theoretical gap. Although the Möller-Nix formula seems to handle unpaired protons as a function of shape relatively well, a systematic underprediction of the neutron pairing ener-



FIG. 6. Mass formulae comparisons for krypton and rubidium isotopes with N = 37 - 43. All values are taken from Ref. 27.

gy seems evident in the krypton isotopes. A systematic study of the strontium masses would help test this hypothesis. The special case of the selfconjugate nucleus, ⁷⁶Sr, would permit testing of the Wigner energy terms in various mass models. Thus, even though the present study adds two pieces to the nuclidic mass surface between the $f_{7/2}$ and $g_{9/2}$ shells, it is clear that remeasurements and new values must be obtained. These mass data can then be combined with nuclear structure information to also provide tests for the degree of shape changes in this portion of the chart of the nuclides.

ACKNOWLEDGMENTS

We wish to thank Ms. C. J. Woodward for her assistance with the data analysis. Oak Ridge National Laboratory is operated by Union Carbide Corporation for the U.S. Department of Energy under Contract No. W-7405-eng-26. The work at Texas A & M was also supported in part by the U. S. Department of Energy and the R. A. Welch Foundation.

- ¹J. C. Hardy, J. A. Macdonald, H. Schmeing, T. Faestermann, H. R. Andrews, J. S. Geiger, R. L. Graham, and K. P. Jackson, Phys. Lett. <u>63B</u>, 27 (1976).
- ²J. A. Macdonald, J. C. Hardy, H. Schmeing, T. Faestermann, H. R. Andrews, J. S. Geiger, R. L. Graham, and K. P. Jackson, Nucl. Phys. <u>A288</u>, 1 (1977).
- ³J. C. Hardy, T. Faestermann, H. Schmeing, J. A. Macdonald, H. R. Andrews, J. S. Geiger, R. L. Graham, and K. P. Jackson, Nucl. Phys. <u>A371</u>, 349 (1981).
- ⁴M. Epherre, G. Audi, C. Thibault, R. Klapisch, G. Huber, F. Touchard, and H. Wollnik, Phys. Rev. C <u>19</u>, 1504 (1979).
- ⁵G. Audi, M. Epherre, C. Thibault, A. H. Wapstra, and K. Bos, Nucl. Phys. <u>A378</u>, 443 (1982).
- ⁶D. M. Moltz, K. S. Toth, F. T. Avignone, III, H. Noma, B. G. Ritchie, and B. D. Kern, Phys. Lett. <u>113B</u>, 16 (1982).
- ⁷T. A. Girard and F. T. Avignone, III, Nucl. Instrum. Methods <u>154</u>, 199 (1978).
- ⁸F. T. Avignone, III, L. P. Hopkins, and Z. D. Greenwood, Nucl. Sci. Eng. <u>72</u>, 216 (1979).
- ⁹F. T. Avignone, III, H. Noma, D. M. Moltz, and K. S. Toth, Nucl. Instrum. Methods <u>189</u>, 453 (1981).
- ¹⁰H. Schmeing, J. C. Hardy, R. L. Graham, and J. S. Geiger, Nucl. Phys. <u>A242</u>, 232 (1975).
- ¹¹S. Matuski, N. Sakamoto, K. Ogino, Y. Kadota, T. Tanabe, and Y. Okuna, Nucl. Phys. <u>A370</u>, 1 (1981).
- ¹²R. E. Tribble, J. D. Cossairt, D. P. May, and R. A. Kenefick, Phys. Rev. C <u>16</u>, 917 (1977).
- ¹³R. E. Tribble, R. A. Kenefick, and R. L. Spross, Phys.

Rev. C 13, 50 (1976).

- ¹⁴R. E. Tribble, J. D. Cossairt, and R. A. Kenefick, Phys. Lett. <u>61B</u>, 353 (1976).
- ¹⁵R. E. Tribble, D. M. Tanner, and A. F. Zeller, Phys. Rev. C <u>22</u>, 17 (1980).
- ¹⁶R. E. Tribble, J. D. Cossairt, K-I. Kubo, and D. P. May, Phys. Rev. Lett. <u>40</u>, 13 (1978).
- ¹⁷A. H. Wapstra and K. Bos, At. Data Nucl. Data Tables <u>19</u>, 175 (1977).
- ¹⁸R. E. Tribble et al. (unpublished).
- ¹⁹S. Jang and B. Sørensen, Nucl. Phys. <u>A344</u>, 315 (1980).
- ²⁰M. Hamm, C. W. Towsley, R. Hanus, K. G. Nair, and K. Nagatani, Phys. Rev. Lett. <u>36</u>, 846 (1976).
- ²¹D. H. Lueders, J. M. Daley, S. G. Buccino, F. E. Durham, C. E. Hollandsworth, W. P. Bucher, and H. D. Jones, Phys. Rev. C 11, 1470 (1975).
- ²²C. J. Lister, P. E. Haustein, D. E. Alburger, and J. W. Olness, Phys. Rev. C <u>24</u>, 260 (1981).
- ²³G. K. Bavaria, J. E. Crawford, S. Calamawy, and J. E. Kitching, Z. Phys. A <u>302</u>, 329 (1981).
- ²⁴G. T. Garvey and I. Kelson, Phys. Rev. Lett. <u>16</u>, 197 (1966).
- ²⁵G. T. Garvey, W. J. Gerace, R. L. Jaffe, I. Talmi, and I. Kelson, Rev. Mod. Phys. <u>41</u>, S1 (1969).
- ²⁶J. Jänecke, At. Data Nucl. Data Tables <u>17</u>, 455 (1976).
- ²⁷S. Maripuu and K. Way, At. Data Nucl. Data Tables <u>17</u>, 410 (1976).
- ²⁸P. Möller and J. R. Nix, At. Data Nucl. Data Tables <u>26</u>, 165 (1981).
- ²⁹P. Möller and J. R. Nix, Nucl. Phys. <u>A361</u>, 117 (1981).